

AMENDMENTS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Twice Amended) Process for the preparation of a retarded release formulation for oral use in tablet form, containing at least one active principle with a pharmaceutical, dietary or alimentary action in amounts of between 70-95% by weight, and at least one hydrogenated fatty acid, as the only vehicle, in amounts of between 5 and 30%, relative to the weight of the formulation, wherein:
 - (a) said at least one active principle is mixed with said at least one hydrogenated fatty acid in the melted state in the weight proportions defined above without the addition of any excipients and/or adjuvants;
 - (b) the blend thus obtained is cooled to 5-20 °C and then granulated using a granulator having holes with a diameter of between 1 and 4 mm;
 - (c) the granules thus obtained are then compressed.
2. (Original) The process according to claim 1, wherein said hydrogenated fatty acid is present in amounts between 10 and 20%, relative to the weight of the formulation.
3. (Original) The process according to claim 1, wherein the blend in point (b) is cooled to 10⁰C -12⁰C.

4. (Original) The process according to claim 1, wherein the blend in point (b) is granulated using a granulator having holes with a diameter of between 1 and 2 mm.
5. (Previously Amended) The process according to claim 1 wherein: before the mixing in point (a), said at least one active principle is premixed at ambient temperature with excipients and/or adjuvants.
6. (Previously Amended) A formulation obtainable by the process according to any of claims 1-5.
7. (Previously Amended) A formulation according to claim 6, characterized in that said at least one hydrogenated fatty acid has a chain comprising between 3 and 20 carbon atoms, or mixtures thereof.
8. (Original) A formulation according to claim 6, characterized in that said at least one hydrogenated fatty acid is hydrogenated palm oil.
9. (Twice Amended) A formulation obtainable by the process according to any of claims 1-4 [according to claim 6], characterized in that said at least one active principle is present in an amount of 70-95%, relative to the weight of the formulation, and in that said at least one active principle and said at least one hydrogenated fatty acid make up 100% by weight of the formulation.

10. (Previously Amended) A formulation according to claim 6, characterized by containing:
- (a) from 10 to 50% by weight, of at least one active principle with a pharmaceutical, dietary or alimentary action; (b) from 5 to 30% by weight of said fatty acid and (c) excipients and/or adjuvants, the sum of the components (a), (b) and (c) making up 100% by weight of the formulation.
11. (Original) A formulation according to claim 10, characterized by containing from 30 to 50% by weight of component (a).
12. (Original) A formulation according to claim 10, characterized by containing from 20 to 30% by weight of component (b).
13. (Original) A formulation according to claim 6, characterized in that said at least one active principle with a therapeutic action is selected from non-steroid and steroid anti-inflammatory drugs, tranquilizers, sleeping pills, anti-hypertensive, anti-histaminic and anti-asthmatic drugs and in that said at least one active principle with a dietary or alimentary action is selected from the group consisting of lactic acid microorganisms, beer yeasts, either as such or containing living cells, vitamins, minerals, amino acids, vegetable extracts, and derivatives thereof.
14. (Previously Added) The process according to claim 1, wherein the blend in point (b) thus obtained is cooled to 10°C -12°C.

15. (Previously Added) The process according to claim 1, wherein the blend in point (b) thus obtained is granulated after cooling using a granulator having holes with a diameter of between 1 and 2 mm.

16. (Previously Added) A formulation according to claim 6, characterized in that said at least one hydrogenated fatty acid has a chain comprising between 8 and 14 carbon atoms, or mixtures thereof.

17. (Currently Amended) A formulation obtainable by the process according to any of claims 1-4 [according to claim 6], characterized in that said at least one active principle is present in an amount of 75-90%, relative to the weight of the formulation, and in that said at least one active principle and said at least one hydrogenated fatty acid make up 100% by weight of the formulation.

18. (New) A formulation according to claim 1, characterized in that said at least one active principle is present in an amount of 75-90%, relative to the weight of the formulation.